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=> s n-glycoylneuraminic acid
             6 N-GLYCOYLNEURAMINIC ACID
L1
=> s n-glycolylneuraminic acid
          1191 N-GLYCOLYLNEURAMINIC ACID
L2
=> s (11 and 12) and (retrovirus or hiv or aids)
             0 (L1 AND L2) AND (RETROVIRUS OR HIV OR AIDS)
L3
=> s (11 or 12) and (retrovirus or hiv or aids)
             6 (L1 OR L2) AND (RETROVIRUS OR HIV OR AIDS)
L4
=> dup rem
ENTER L# LIST OR (END):14
PROCESSING COMPLETED FOR L4
              2 DUP REM L4 (4 DUPLICATES REMOVED)
=> d 15 1-2 bib ab
                                                         DUPLICATE 1
     ANSWER 1 OF 2 MEDLINE
T.5
     94148878
                  MEDLINE
ΑN
                PubMed ID: 8106417
     94148878
DN
     Generation of Chinese hamster ovary cell glycosylation mutants by
TΙ
     retroviral insertional mutagenesis. Integration into a discrete locus
      generates mutants expressing high levels of N-
      glycolylneuraminic acid.
      Hubbard S C; Walls L; Ruley H E; Muchmore E A
 ΑU
     Center for Cancer Research, Massachusetts Institute of Technology,
 CS
      Cambridge 02139.
      R01 HG00684 (NHGRI)
 NC
      R01-CA40602 (NCI)
      R29-GM43165 (NIGMS)
      JOURNAL OF BIOLOGICAL CHEMISTRY, (1994 Feb 4) 269 (5) 3717-24.
 SO
      Journal code: HIV; 2985121R. ISSN: 0021-9258.
      United States
 CY
      Journal; Article; (JOURNAL ARTICLE)
 DT
      English
 LA
      Priority Journals
 FS
      199403
 EM
      Entered STN: 19940330
 ED
      Last Updated on STN: 19980206
      Entered Medline: 19940318
      Retroviral insertional mutagenesis can both generate somatic cell mutants
 AB
      and pinpoint the genomic locus associated with a mutant phenotype. In the
      present study, this approach was applied to Chinese hamster ovary cells
      (CHO) made susceptible to Moloney murine leukemia virus (MoMuLV)
 infection
      by stable expression of an ecotropic retrovirus receptor. These
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CHO cells were infected with a replication incompetent MoMuLV construct with a promoterless hygromycin phosphotransferase (hygro) gene inserted into the U3 region of the long terminal repeat and a second selectable marker, neomycin phosphotransferase (neo), expressed from an internal promoter. CHO clones containing integrated proviruses were selected with hygromycin or G418, and the subset of these with reduced cell surface Neu5Ac were then selected with wheat germ agglutinin (WGA). The majority of the resulting clones had a phenotype not previously described for WGA-resistant CHO mutants arising spontaneously or from chemical mutagenesis: Neu5Ac was almost completely replaced by Neu5Gc. We have provisionally termed these clones SAP mutants, for sialic acid phenotype. Southern analysis of HindIII digested DNA from four SAP mutants revealed that the MoMuLV provirus is present in a 10.4-kilobase (kb) fragment. Probing with a flanking CHO sequence resulted in equivalent hybridization to a 4.6-kb fragment and the 10.4-kb provirus-containing fragment in all four cases, while uninfected parental cells and non-SAP glycosylation mutants generated in the same retrovirus insertional mutagenesis experiments yielded only the 4.6-kb fragment. Sequencing of the 3'-flanking DNA revealed that each of the four SAP mutants had a unique provirus integration site falling within a 796 bp region of the CHO genome. The frequency with which SAP mutants arise suggests that this may be a preferred site for retrovirus integration.

- ANSWER 2 OF 2 BIOSIS COPYRIGHT 2001 BIOSIS L5
- 1994:47135 BIOSIS AN
- PREV199497060135 DN
- Retrovirus insertional mutagenesis of an 796 bp locus in the CHO ΤI genome generates mutants expressing high levels of ${ t N}$ glycolylneuraminic acid.
- Hubbard, S. Catherine (1); Walls, Lorraine; Ruley, H. Earl; Muchmore, ΑU Elaine A.
- (1) Genzyme Corp., One Kendall Square, Cambridge, MA 02139 USA CS
- Glycobiology, (1993) Vol. 3, No. 5, pp. 534. SO Meeting Info.: 22nd Annual Meeting of the Society for Complex Carbohydrates San Juan, Puerto Rico November 17-20, 1993 ISSN: 0959-6658.
- Conference DT
- LА English